

**PROFESSIONAL SOLUTIONS - HAND SANITIZER FOAM- benzalkonium chloride liquid
Cosmetic Solutions LLC**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Professional Solutions® - Hand Sanitizer Foam

DRUG FACTS:

ACTIVE INGREDIENTS

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

USES

To decrease bacteria on the skin

WARNINGS

For external use only: Hands only

When using this product avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. Avoid contact with broken skin.

Stop use and seek a doctor if redness or irritation develops and persist for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

Wet hands thoroughly with product and allow to dry without wiping - for children under 6, use only under adult supervision - not recommended for infants.

INACTIVE INGREDIENTS

Water (Aqua), Propanediol, Glycereth-7 Triacetate, Glycerin, Babassu Oil Glycereth-8 Esters, Decyl Glucoside, Calendula Officinalis Flower Extract, Equisetum Arvense Extract, Geranium Maculatum Extract, Honey, Citric Acid, Panax Ginseng Root Extract, Salvia Officinalis Leaf Extract, Sambucus Nigra Flower Extract, Ethylhexylglycerin, Phenoxyethanol, Sodium Hydroxide.

Manufactured By: Professional Solutions
Boca Raton, FL 33487

PRINCIPAL DISPLAY PANEL - 50 mL Bottle Label

Professional
Solutions®

PURE SCIENCE. BETTER SKIN

hand sanitizer
foam
protect + moisturize
1.7 fl. oz. (50mL)



Professional Solutions
PURE SCIENCE. BETTER SKIN

hand sanitizer foam

protect + moisturize

1.7 fl. oz. (50mL)

Manufactured By: Professional Solutions
Boca Raton, FL 33487
www.professionalsolutions.net - Made in USA



hand sanitizer foam

DRUG FACTS:	
ACTIVE INGREDIENTS: Benzalkonium Chloride 0.13%	Purpose: Antiseptic
USES: To decrease bacteria on the skin	
WARNINGS:	
For external use only: Hands only	
When using this product avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. Avoid contact with broken skin.	
Stop use and seek a doctor if redness or irritation develops and persist for more than 72 hours.	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
DIRECTIONS: Wet hands thoroughly with product and allow to dry without wiping - for children under 6, use only under adult supervision - not recommended for infants.	
INACTIVE INGREDIENTS: Water (Aqua), Propenediol, Glycereth-7 Triacetate, Glycerin, Babassu Oil, Glycereth-8 Esters, Decyl Glucoside, Calendula Officinalis Flower Extract, Equisetum Arvense Extract, Geranium Maculatum Extract, Honey, Citric Acid, Panax Ginseng Root Extract, Salvia Officinalis Leaf Extract, Sambucus Nigra Flower Extract, Ethylhexylglycerin, Phenoxyethanol, Sodium Hydroxide.	



PROFESSIONAL SOLUTIONS - HAND SANITIZER FOAM

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66163-1900
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	

Propanediol (UNII: 5965N8W85T)	
Glycereth-7 Triacetate (UNII: S9W6Z48HUP)	
Glycerin (UNII: PDC6A3C0OX)	
Babassu Oil Glycereth-8 Esters (UNII: 12XQ76K0YL)	
Decyl Glucoside (UNII: Z17H97EA6Y)	
Calendula Officinalis Flower (UNII: P0M7O4Y7YD)	
Equisetum Arvense Branch (UNII: 1L0VKZ185E)	
Geranium Maculatum Root (UNII: 93IXI5B6OJ)	
Honey (UNII: Y9H1V576FH)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Asian Ginseng (UNII: CUQ3A77YXI)	
Sambucus Nigra Flower (UNII: 07V4DX094T)	
Ethylhexylglycerin (UNII: 147D247K3P)	
Phenoxyethanol (UNII: HIE492ZZ3T)	
Sodium Hydroxide (UNII: 55X04QC32I)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66163-1900-0	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	
2	NDC:66163-1900-1	120 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	05/01/2020	

Labeler - Cosmetic Solutions LLC (807907928)